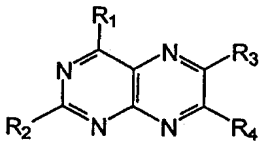




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(57) Abstract The invention relates to a pharmaceutical composition for the treatment of autoimmune disorders or for the treatment and/or prevention of transplant-rejections and/or for the treatment of inflammatory diseases comprising as an active principle at least one pteridine derivative having general formula (I). The invention further relates to a compound having said general formula and to the use of a compound as a medicament and for the manufacture of a medicament for the treatment of autoimmune disorders, for the prevention of transplant-rejections and for the treatment of inflammatory diseases.		

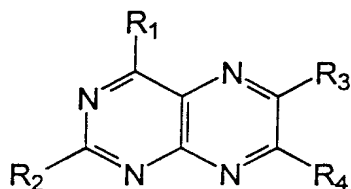
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IMMUNOSUPPRESSIVE EFFECTS OF PTERIDINE DERIVATIVES

The invention relates to a pharmaceutical composition for the treatment of autoimmune disorders or for the treatment and/or prevention of transplant-rejections and/or the treatment of inflammatory diseases comprising as active ingredient one or more pteridine derivatives having the general formula:



(I)

wherein:

R₁ and R₂ are independently amino, hydroxylamino, alkoxyamino, hydrazino, piperazino, N-alkylpiperazino, morpholino, mono- and diarylamino, (wherein the aryl group may be the same or different) mono- and dialkylamino (wherein the alkyl group may be the same or different), mono- and diarylalkylamino (wherein both groups may be the same or different), cycloalkylamino (such as cyclopropylamino, cyclobutylamino, cyclopentylamino, cyclohexylamino), alkoxy, mercaptoalkyl. The alkyl group may contain 1 to 7 carbon atoms and may be branched, cyclized and may be oxidized;

R₃ : unsubstituted, monosubstituted or disubstituted aryl group (wherein the substituent may be, but not limited to, halogen, alkoxy, alkyl), aryl group bond to the pteridine ring via a saturated or unsaturated alifatic spacer which may be halogenated or hydroxylated, aliphatic substituent which may contain ether function, alcohol function, substituted or unsubstituted amino functions; and

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R₄ : hydrogen, alkyl, alkoxy, substituted or unsubstituted aryl.

The invention further relates to combined pharmaceutical preparations comprising one or more pteridine derivates and one or more known immunosuppressant, and to a group of novel pteridine derivates as such.

Further the invention is also related to a method for the treatment of autoimmune disorders and/or of transplant-rejections and/or inflammatory diseases.

The invention further relates to a method for the preparation of the above mentioned pteridine derivatives and the the pteridine derivatives as such.

Several pteridine derivates are known in nature and used in the preparation of medicines, for example as described in EP-A-108 890. Other medical uses of derivates of pteridine are described in WO 95-31987 as NO-synthase inhibitors for example for the treatment of diseases caused by a high nitrogen monoxide level. Further, WO-95-32203 describes also the use of tetrahydropteridine derivates as NO-synthase inhibitors.

Both above-mentioned WO publications disclose also the use of specific pteridine derivates in the treatment of pathologically low blood pressure, in particular septic shock and combined with cytokines in tumor therapy and in transplant-rejection diseases.

Although some of these pteridine derivates are claimed as potentially active for the treatment of transplant-rejection diseases, direct evidence for their effectiveness is lacking. Overall there still is a need for specific and highly active immunosuppressive compounds, in particular immunosuppressive compounds active in the cosignal pathway.

A first object of the invention is to provide a pharmaceutical composition having high immunosuppressive activity. Another object of the invention is to provide a

combined immunosuppressive preparation which causes a superadditive effect, comprising a pteridine derivate of the invention and other known immunosuppressants.

Another further object of the invention is to provide immunosuppressive compounds, which are active in a minor dose, in order to decrease the considerable treatment costs.

Known immunosuppressive compounds are for example cyclosporine A, substituted xanthines, tacrolimus (FK 506), rapamycin (RPM), leflunomide, mofetil, adrenocortical steroids, cytotoxic drugs and antibody preparations.

The immunosuppressive effect of cyclosporine A (CyA) is already known since 1972. However, due to its nephrotoxicity and several other side effects CyA has not been able to establish itself as the optimal and final drug of choice.

Methylxanthines, for example pentoxifylline (PTX), are known having immunosuppressive effects in vitro.

Recently (Lin Y. et al, Transplantation 63 (1997) it has been found that the co-medication of an immunosuppressive compound such as cyclosporine A (CyA) or FK506 or RPM (rapamycin) with a methylxanthine derivative, in particular A802715 (7-propyl-1(5-hydroxy-5-methylhexyl)-3-methylxanthine) leads to a superadditive increase in the immunosuppressive action.

Likewise, other substituted, in particular substituted 8-phenylxanthines have been found to possess immunosuppressive effects in vitro (application EP 98.201323.7).

The present invention relates in particular to the application of a group pteridine derivatives and their pharmaceutical salts, possessing unexpectedly desirable pharmaceutical properties, i.e. are highly active immunosuppressive agents, are useful in the treatment in

transplant rejection and/or in the treatment of inflammatory diseases.

The invention demonstrates the immunosuppressive effects of pharmaceutical compositions for the treatment of autoimmuno disorders or of transplant-rejections comprising one or more pteridine derivatives of the above formula (I) or salts thereof.

The term pharmaceutically acceptable addition salt as used hereinbefore defines the non-toxic, therapeutically active addition salt forms which the compounds of formula (I) may form. The compounds of formula (I) having basic properties may be converted into the corresponding therapeutically active, non-toxic acid addition salt forms by treating the free base form with a suitable amount of an appropriate acid following conventional procedures. Examples of appropriate acids are for example, inorganic acids, for example, hydrohalic acid, e.g. hydrochloric, hydrobromic and the like acids, sulfuric acid, nitric acid, phosphoric acid and the like; or organic acids, such as, for example, acetic, propanic, hydroxyacetic, 2-hydroxypropanic, 2-oxopropanic, ethanedioic, propanedioic, butanedioic, (Z)-2-butenedioic, (E)-2-butenedioic, 2-hydroxybutanedioic, 2,3-dihydroxybutanedioic, 2-hydroxy-1,2,3-propanetricarboxylic, methanesulfonic, ethanesulfonic, benzenesulfonic, 4-methylbenzenesulfonic, cyclohexanesulfamic, 2-hydroxybenzoic, 4-amino-2-hydroxybenzoic and the like acids.

The compounds of formula (I) having acidic properties may be converted in a similar manner into the corresponding therapeutically active, non-toxic base addition salt forms. Examples of such base addition salt forms are, for example, the sodium, potassium, calcium salts, and also the salts with pharmaceutically acceptable amines such as, for example, ammonia, alkylamines, benzathine, N-methyl-D-glucamine, hydrabamine, amino acids, e.g. arginine, lysine. The term pharmaceutically

acceptable addition salts also comprises the solvates which the compounds of formula (I) may form, e.g. the hydrates, alcoholates and the like.

The term stereochemically isomeric forms as used hereinbefore defines the possible different isomeric as well as conformational forms which the compounds of formula (I) may possess. Unless otherwise mentioned or indicated, the chemical designation of compounds denotes the mixture of all possible stereochemically and conformationally isomeric forms, said mixtures containing all diastereomers, enantiomers and/or conformers of the basic molecular structure. All stereochemically isomeric forms of the compounds of formula (I) both in pure form or in admixture with each other are intended to be embraced within the scope of the present invention.

Some compounds of the present invention may exist in different tautomeric forms and all such tautomeric forms are intended to be included within the scope of the present invention.

The compounds of the present invention show a broad spectrum profile as is evidenced by the results obtained in the diversity of test procedures cited hereinbefore.

An advantageous feature of the compounds of the present invention resides in their excellent oral activity; the present compounds when administered orally have been found to be practically equipotent with the same being administered subcutaneously.

A particularly important asset of most of the present compounds is their lack of sedating properties at therapeutic dose levels, a troublesome side effect associated with many antihistaminic and antiallergic compounds. The non-sedating properties of the present compounds can be demonstrated, for example, by the results obtained in studying the sleep - wakefulness cycle of the rat (Psychopharmacology, 97, 436-442, (1989)).

Another interesting feature of the present compounds relates to their fast onset of action and the favorable duration of their action.

In view of their useful properties the subject compounds may be formulated in various pharmaceutical forms for administration purposes. To prepare the antiallergic compositions of this invention, an effective amount of the particular compound, in base or acid addition salt form, as the active ingredient is combined in intimate admixture with a pharmaceutically acceptable carrier, which carrier may take a wide variety of forms depending on the form of preparation desired for administration. These pharmaceutical compositions are desirably in unitary dosage form suitable, preferably, for administration orally, rectally, percutaneously, or by parenteral injection. For example, in preparing the compositions in oral dosage form, any of the usual pharmaceutical media may be employed such as, for example, water, glycols, oils, alcohols and the like in the case of oral liquid preparations such as suspensions, syrups, elixirs and solutions; or solid carriers such as starches, sugars, kaolin, lubricants, binders, disintegrating agents and the like in the case of powders, pills, capsules and tablets. Because of their ease in administration, tablets and capsules represent the most advantageous oral dosage unit form, in which case solid pharmaceutical carriers are obviously employed. For parental compositions, the carrier will usually comprise sterile water, at least in large part, through other ingredients, for example to aid solubility, may be included. Injectable solutions, for example, may be prepared in which the carrier comprises saline solution, glucose solution or a mixture of saline and glucose solution. Injectable suspensions may also be prepared in which case appropriate liquid carriers, suspending agents and the like may be employed. In the compositions suitable for percutaneous administration, the

carrier optionally comprises a penetration enhancing agent and/or a suitable wetting agent, optionally combined with suitable additives of any nature in minor proportions, which additives do not introduce a significant deleterious effect on the skin. Said additives may facilitate the administration to the skin and/or may be helpful for preparing the desired compositions. These compositions may be administered in various ways, e.g. as a transdermal patch, as a spot-on or as an ointment. Acid addition salts of the subject compounds due to their increased water solubility over the corresponding base form, are obviously more suitable in the preparation of aqueous compositions.

It is especially advantageous to formulate the aforementioned pharmaceutical compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form as used in the specification and claims herein refers to physically discrete units suitable as unitary dosages, each unit containing a predetermined quantity of active ingredient calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier. Examples of such dosage unit forms are tablets (including scored or coated tablets), capsules, pills, powder packets, wafers, injectable solutions or suspensions, teaspoonfuls, tablespoonfuls and the like, and segregated multiples thereof.

The present invention also relates to a method of treating warm-blooded animals suffering from said allergic diseases by administering to said warm-blooded animals an effective antiallergic amount of a compound of formula (I).

In general it is contemplated that an effective antiallergic amount would be from about 0.001 mg/kg to about 20 mg/kg body weight, and more preferably from about 0.01 mg/kg to about 5 mg/kg body weight.

The following examples are intended to illustrate and not to limit the scope of the present invention.

only in all its aspects.

Experimental Part

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2-amino-4-n-pentyloxy-6-styrylpteridine (1).

A mixture of 2-amino-6-chloro-4-n-pentyloxypteridine [1] (1.5 g, 5.6 mmols) palladium acetate (63 mg, 0.28 mmols), tri-*o*-tolylphosphane (682 mg, 2.24 mmols),
10 cuprous iodide (53 mg, 0.28 mmols), styrene (1,3 ml., 11.3 mmols) and triethylamine 3.1 ml, 22 mmols) was stirred in dry acetonitrile (50 ml.) under reflux for 90 hours. It was evaporated and the residue purified by silica gel column chromatography with CHCl₃. The product
15 fraction was evaporated to give 1.37 g (72%) of an orange powder. Recrystallization from EtOAc/hexane. M.p. 127-128°C.

2-Amino-6-(1,2-dibromophenethyl)-4-n-pentyloxypteridine (2).

To a solution of **1** (1.0 g, 2.94 mmols) in chloroform (50 ml.) was added a 2 M bromine solution in chloroform (2.2 ml., 4.4 mmols) and then the mixture stirred at room temperature for 7 hours. It was diluted with chloroform
25 (50 ml.), washed with a saturated aqueous Na₂SO₃ solution (100 ml.) and dried over Na₂SO₄. It was evaporated, the residue treated with little toluene, filtered, washed with ether and dried in a vacuum desiccator to give 0.84 g (57%) yellow powder.

30

2-Amino-4,7-dimethoxy-6-styrylpteridine (3).

A suspension of **2** (0.3 g, 0.6 mmols) in abs. Methanol (10 ml.) was treated with 1 M methanolic sodium methoxide (3 ml., 3 mmols) and then refluxed for 4 hours. It was
35 diluted with chloroform (100 ml.), washed with saturated aqueous NH₄Cl solution and water and then the solution

dried over Na_2SO_4 . The filtrate was evaporated and the residue purified by silica gel column chromatography in chloroform. The product fraction was evaporated to give 50 mg. (26%) of a yellow powder, M.p. 197-198°C.

5

O⁴-Methyl-biopterin (4).

To a solution of N^2 , 1',2'-O-triacetyl-biopterin (1,0 g; 2.75 mmoles), triphenylphosphane (12,08 g, 4.13 mmoles) and methanol (0.15 ml., 3.7 mmoles) in dry dioxane (30 ml.) was added diisopropyl azodicarboxylate (0.81 g, 4.11 mmoles) and after stirring for 1.5 hours at room temperature evaporated to dryness. The residue was purified by silica gel column chromatography eluting with EtOAc/ CHCl_3 (1:4). The product fraction was evaporated and dried in vacuum to give 0.4 g (38%) of N^2 , 1',2'-O-triacetyl-O⁴-methylbiopterin.

Deacetylation of the reaction product (0.28 g, 0.74 mmoles) was done by stirring in abs. Methanol (20 ml.) and triethylamine (4 ml.) for 24 hours. Evaporation to dryness, treatment of the residue with ether, filtration and drying gave 0.172 g (83%) of **4**. M.p. 160-161°C (Decomp.).

General procedure for the synthesis of 2,4-diamino-6-arylpteridines (5, 7, 8, 9)

A suspension of 2,4,5,6-tetraaminopyrimidine dihydrochloride (2.13 g, 0.01 moles) in methanol (100 ml.) was heated to boiling and then a solution of the arylglyoxalmonoxime (phenylglyoxalmonoxime [2], p-methylphenylglyoxalmonoxime [3], p-methoxyphenylglyoxalmonoxime [4], p-chlorophenylglyoxalmonoxime [5] (0.015 moles) in methanol (20 ml.) added dropwise within 30 min. It was heated under reflux for 2 hours forming a precipitate. After cooling was neutralized by conc. ammonia to pH 8 with stirring. The precipitate was collected, washed with

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methanol and ether and dried in the oven at 100°C.
Yield: 85-95%. The reaction product is usually
chromatographically pure. Recrystallization can be
achieved from DMF.

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2,4-Diamino-7-methyl-6-phenylpteridine (6).

Analogous to the preceding procedure using α -
hydroximinopropiophenon. Yield: 70%.

10 General procedures for the synthesis of 4-amino-6-aryl-2-
 β -hydroxyethylaminopteridines (10, 11, 12)

A suspension of 4,5,6-triamino-2- β -
hydroxyethylaminopyrimidine trihydrochloride (2.93 g,
0.01 moles) in methanol (60 ml.) was heated under reflux
15 and then a solution of the aryl-glyoxalmonoxime (0.015
moles) in methanol (15 ml.) added dropwise. After reflux
for 2 hours and cooling was neutralized to pH 9 with
conc. Ammonia to give a yellow precipitate. Yield: 90%.

20 2-amino-4-hydroxylamino-6-phenylpteridines (13).

A suspension of 2,5,6-triamino-4-methoxypyrimidine
dihydrochloride (1 g, 4 mmoles) in methanol (40 ml.) was
heated to boiling and then a solution of
phenylglyoxalmonoxime (1 g, 6.6 mmoles) in methanol (10
25 ml.) added dropwise. A clear solution is obtained from
which on reflux for 2 hours a precipitate separated out.
The solid was filtered off (hydrochloride salt),
suspended in water (30 ml) and then neutralized to pH 8
by conc. ammonia. The precipitate was collected, washed
30 with water and ethanol and dried at 100°C to give a
yellow powder. Yield: 0.84 g (82%).

2,4-diamino-6-bromomethylpteridine [6].

A suspension of 2,4,5,6-tetraaminopyrimidine
35 trihydrobromide (3.0 g, 0.01 moles) in methanol (60 ml)
was heated to reflux and then a solution of β -

11
bromopyruivaldoxime (0.015 moles) in methanol (30 ml) added dropwise within 10 min. The resulting yellow solution was refluxed for 30 min., then cooled to room temperature and neutralized by conc. ammonia to pH 8.
5 The yellow precipitate was collected, washed with little methanol and ether and dried in a vacuum desiccator. Yield: 88%.

10 General procedure for 2,4-diamino-6-alkoxymethyl-(17,18) and -6-aminomethylpteridines (19,20).

To a mixture of dimethylacetamide (DMA) (30 ml) and the appropriate alcohol (β -methoxyethanol, n-decanol) (5 ml) was added sodium hydride (1 g, 80%) and after stirring for 1 hour 2,4-diamino-6-bromomethylpteridine (1 g)
15 added. Stirring was continued at room temperature for 6 hours, then diluted with H₂O (100 ml) and kept in the icebox for 2 days. The precipitate was collected and recrystallized from EtOH/ conc. NH₃ (16:1). Yield: 50%.

An analogous reaction takes place with amines
20 (dimethylamine in ethanol, benzylamine) (0.04 mmoles) in DMA (20 ml) and 2,4-diamino-6-bromomethylpteridine (2.55 g, 0.01 moles). Yield: 50-60%.

25 General procedures for the synthesis of 2,6-diamino-4-dialkylamino-5-p-chlorophenylazopyrimidines.

A solution of 2,6-diamino-4-dialkylamino-5-p-chlorophenylazopyrimidine [7] (5.0 g, 16.6 mmoles) in DMF (50ml) and the appropriate amine (dimethylamine in ethanol (50%), diethylamine, di-n-propylamine,
30 dibenzylamine, morpholine, piperidine, pyrrolidine, piperazine, N-methylpiperazine) (10.0 g) was heated in an oilbath to 70°C for 5 hours. Then water (50 ml) was added, cooled and the yellow precipitate collected, washed with water and dried. Recrystallization from EtOH
35 or DMF/water. Yield: 55-90%.

General procedure for the synthesis of 2,5,6-triamino-4-dialkylaminopyrimidines.

A suspension of 2,6-diamino-4-dialkylamino-5-p-chlorophenylazopyrimidine (3.28 g, 10 mmoles) in methanol (70 ml) and conc. ammonia (10 ml) was reduced in a shaking apparatus under H₂ atmosphere in presence of Raney nickel catalyst (3.5 g) for 2 days. The catalyst was filtered off under argon atmosphere and then the filtrate evaporated in vacuo to dryness. The residue was treated with ether to remove the p-chloroaniline, filtered and then the solid stirred in methanolic HCl (10%, 50 ml) overnight. The dihydrochloride salt was collected and dried in a vacuum desiccator over KOH. Yield: 85-90%.

General procedure for the synthesis of 2-amino-4-dialkylamino-6-arylpteridines (14-16, 21-49)

To a boiling solution of the 2,5,6-triamino-4-dialkylaminopyrimidine dihydrochloride salt (5 mmoles) in MeOH (20 ml) was added a solution of the arylglyoxalmonoxime (7.5 mmoles) in MeOH (10 ml) dropwise and then the mixture heated under reflux for 3 hours. After cooling the suspension or solution was made alkaline by conc. ammonia to pH 9 and the resulting precipitate filtered off, washed with water and dried. Recrystallization was done from EtOH and DMF/H₂O, respectively, to give a yellow solid. Yield: 50-85%.

Further compounds 50-66 were prepared according to the above described syntheses and tested.

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Materials and methods

Various models may be used for testing an immunosuppressive effect. In vivo, for example, different transplantation models are available. They are strongly influenced by different immunogenicities, depending on the donor and recipient species used and depending on the nature of the transplanted organ. The survival time of transplanted organs can thus be used to measure the suppression of the immune response. In vitro, there exist also various models. The most used are lymphocyte activation tests. Usually activation is measured via lymphocyte proliferation. Inhibition of proliferation thus always means immunosuppression under the experimental conditions applied. There exist different stimuli for lymphocyte activation:

- coculture of lymphocytes of different species (MLR = mixed lymphocyte reaction): lymphocytes expressing different minor and major antigens of the HLA-DR type (= alloantigens) activate each other non-specifically.
- CD3 assay: here there is an activation of the T-lymphocytes via an exogenously added antibody (OKT3). This antibody reacts against the CD3 molecule located on the lymphocyte membrane. This molecule has a costimulatory function. The interaction anti-CD3 (= OKT3)-CD3 results in T-cell activation which proceeds via the Ca^{2+} /calmodulin/calcineurin system and can be inhibited by CyA.
- CD28 assay: here specific activation of the T-lymphocyte goes also via an exogenously added antibody against the CD28 molecule. This molecule is also located on the lymphocyte membrane, and delivers strong costimulatory signals. This activation is Ca^{2+} -independent and thus cannot be inhibited by CyA.

35 Reagents

All derivatives were dissolved in 0.5 ml DMSO and

15

further diluted in culture medium before use in in vitro experiments. The culture medium consisted of RPMI-1640 + 10% FCS.

5 Mixed Lymphocyte Reaction

Peripheral blood mononuclear cells (PBMC) were isolated from heparinized peripheral blood by density gradient centrifugation over Lymphoprep (Nycomed, Maorstua, Norway). Allogeneic PBMC or EBV-transformed
10 human B cells [RPMI1788 (ATCC name CCL156)] which strongly express B7-1 and B7-2 were used as stimulator cells after irradiation with 30 Gy. MLR was performed in triplicate wells. After 5 days incubation at 37EC, 1 μ Ci [³H]-thymidine was added to each cup. After a further 16
15 hours incubation, cells were harvested and counted in a β -counter.

The percent suppression of proliferation by drugs was counted using the formula

$$\text{Percent inhibition} = \frac{(\text{cpm} + \text{drugs}) - \text{cpm Cult. Med}}{(\text{cpm} - \text{drugs}) - \text{cpm Cult. Med.}} \times 100$$

20

T cell purification

T cells were purified by removing non-T cells. Briefly, monocytes were removed by cold agglutination. The
25 resulting lymphoid cells were further purified by a cell enrichment immunocolumn [Collect Human T (Biotex, Edmonton, Alberta, Canada)] by a process of negative selection. More than 95% of the B cells were removed with this procedure. After depletion, the resulting T cell
30 preparation was highly purified explaining these cells could not be activated by PHA or rIL-2 alone at concentrations capable of stimulating RBMC prior to deletion.

35 Measurements of T cell proliferations induced by anti-CD3

16

mAb + PMA or anti-CD28 mAb + PMA

Highly purified T cells ($10^6/\text{ml}$) were stimulated by immobilized anti-CD3 or anti-CD28 mAb in the presence of PMA. Anti-CD3 mAb (CLB-CD3; CLB, Amsterdam, The Netherlands) were fixed on the 96-microwell plates by incubating the wells with 50 μl of mAb solution (1/800 dilution in culture medium). Anti-CD28 mAb (CLB-CD28; CLB, Amsterdam, The Netherlands) 50 μl (1/650 dilution in culture medium) was added directly to the wells. Further, 20 μl PMA (Sigma, St. Louis, MO, USA) solution (final concentration: 0.5 ng/ml) was added. Subsequently, 20 μl of immunosuppressants were added by serial dilution in triplicate wells. Finally 100 μl of the T cell suspension ($10^6/\text{ml}$) was added. After 48-hour incubation at 37°C in 5% CO_2 20 ϕl BrdU (100 ϕM solution) (Cell Proliferation Elisa, Boehringer-Mannheim Belgium) was added to each well. After a further overnight incubation the T cell proliferation was measured using a colorimetric immunoassay for qualification of cell proliferation based on measurements of the incorporation of BrdU during DNAsynthesis. The optical density (OD) was measured by a Behring EL311 plate reader at 450 nm (reference wavelength: 690 nm). The percent suppression of proliferation by drugs was counted using the formula:

$$\text{Per cent inhibition} = \frac{(\text{OD} + \text{drugs}) - (\text{OD Cult. Med.})}{(\text{OD} - \text{drugs}) - (\text{OD Cult. Med.})} \times 100$$

In vitro immunosuppressive effect of Pteridine Derivatives as measured with the MLR and with tests involving polyclonal T cell proliferation induced by anti-CD3 mAb + PMA or anti-CD28 mAb + PMA (table II)

- Table II shows the IC₅₀ values of the various substances in the MLR. The IC₅₀ value represents the lowest concentration of the substances that resulted in a 50% suppression of the MLR. These concentrations are divided into for subranges i.e.

0 stands for concentrations of at least 151 μ M,
+ stands for concentrations 16-150 μ M,
++ stands for concentrations 1-15 μ M,
+++ stands for concentrations lower than 1 μ M,

10

- Column III shows the IC₅₀ value of the various substances for the anti-CD3 mAb + PMA pathway and row IV the IC₅₀ values of the various substances for the anti-CD28 mAb + PMA pathway.
- As a comparison the values of other immunosuppressants: CsA, FK506, Rapamycin, Leflunomide and Mycophenolic acid methatroxate (MTX) and 5-Fluoro-uracil (5-FU) in table III are given as well.

First, most of the pteridine classes (I) according to the invention contain compounds with a clear suppressive effect in the MLR (mixed lymphocyte reaction). The MLR is considered as an in vitro analogue of the transplant rejection as it is based on the recognition of allogeneic MHC (major histocompatibility antigens) on the stimulator leucocytes, by responding lymphocytes. Various established immunosuppressive drugs are known to suppress the MLR, and were also shown in this description.

From these data it can be deduced that the pteridine derivatives are effective in clinical situations where other immunosuppressants are active as well.

These include the prevention and/or treatment of

organ transplant rejection, the prevention and/or treatment of both rejection and the occurrence of graft-versus-host-disease after BM transplantation; the prevention and/or treatment of autoimmune diseases including diabetes mellitus, multiple sclerosis, glomerulonephritis, rheumatoid arthritis, psoriasis systemic diseases such as vasculitis; scleroderma, polymyositis, autoimmune endocrine disorders (thyroiditis), ocular diseases (uveitis), inflammatory bowel diseases (Crohn's disease, colitis ulcerosa), autoimmune liver diseases (autoimmune hepatitis, primary biliary cirrhosis) autoimmune pneumonitis and auto-immune carditis.

Whereas cyclosporine A and FK506 are only active in the anti-CD3 + PMA test, the pteridine derivatives according to the invention were active, not only in the anti-CD3 + PMA but also in the anti-CD28 + PMA test. It has been shown that the latter is Ca-calmodulin resistant, and resistant to CsA and FK506. The anti-CD28 + PMA pathway has also been called the cosignal pathway and is important to induce energy and even tolerance in T cells. Moreover, representative compounds have been found to be active in an whole blood assay.

Under the term "organ" in the description is understood all organs or parts of organs (even several) in mammals, in particular humans, for example kidney, heart, skin, liver, muscle, cornea, bone, bone marrow, lung, pancreas, intestine or stomach.

After organ transplantation, rejection of the transplanted organ by the recipient occurs (host-versus-graft reaction). After bone marrow transplantation, also rejection of the host by the grafted cell may occur (graft-versus-host reaction). Rejection reactions mean all reactions of the recipient body or of the

transplanted organ which in the end lead to cell or tissue death in the transplanted organ or adversely affect the functional ability and viability of the transplanted organ or adversely affect the functional ability and viability of the transplanted organ or the recipient. In particular, this means acute and chronic rejection reactions.

Auto-immune disorders include, inter alia, systemic lupus erythematosus, rheumatoid arthritis, psoriasis, pemphigus, atopic dermatitis, myositis, multiple sclerosis, nephrotic syndrome (in particular glomerulonephritis), ulcerative colitis or juvenile diabetes.

An additive or synergetic effect of pteridine derivatives and other immunosuppressants may be anticipated. This may be especially, although not exclusively the case for combinations with CyA or FK 506 as the latter are not suppressive in the aCD28 pathway of T cell activation (table III) whereas most Pteridine derivatives are.

The invention further relates to the use of cyclosporin A or FK506 or Rapamycin and at least one pteridine derivative according to the invention for the production of a pharmaceutical for inhibiting the replication of viruses such as picorna-, toga-, bunya-, orthomyxo-, paramyxo-, rhabdo-, retro-, arena-, hepatitis B-, hepatitis C-, hepatitis D-, adeno-, vaccinia-, papilloma-, herpes-, varicella-zoster-virus or human immunodeficiency virus (HIV); or for treating of cancer such as lung cancers, leukaemia, ovarian cancers, sarcoma, Kaposi's sarcoma, meningioma, colon cancers, lymph node tumors, glioblastoma multiforme, prostate cancers or skin carcinomas.

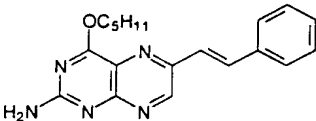
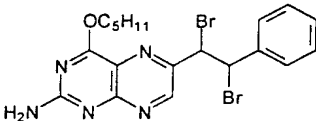
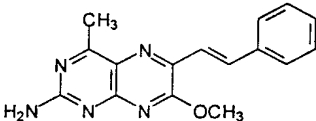
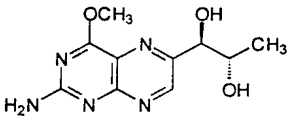
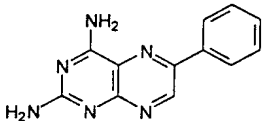
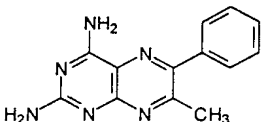
The invention further relates to the use of

cyclosporin A or FK506 or rapamycin and at least one pteridine derivative of the general formula (I) for the production of a pharmaceutical for the treatment of human after organ transplantation or of (auto)immune disorders.

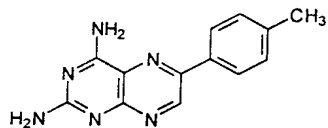
5 Hence, the advantage to associate pteridine with other immunosuppressants may be that, first, the therapeutic spectrum of action of the individual components is quantitatively and qualitatively broadened. Secondly that it allows, by means of a dose reduction
10 without reduced efficacy but with increased safety, that the treatment of immune disorders which were hitherto no indication for immunosuppressive therapy as a result of side effects may be considered. At the same time, the therapy costs can be decreased to an appreciable extent.

15 As a comparison, known pteridine derivatives are submitted to the same test conditions as the pteridine derivatives of the invention. These compounds and the results thereof are given in table IV and show no particular immunosuppressive activity.

20 As been stated above the invention also relates to new pteridine derivatives as such, in particular the compounds 1,2,3,6, 14-16 and 21-66 and their pharmaceutically acceptable salts.

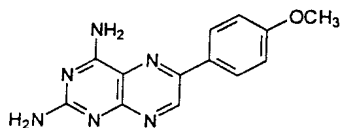
Compound n°	MLR	aCD3	aCD28
1 2-amino-4-pentoxo-6-styrylpteridine			
	15	110	80
2 2-amino-4-n-pentoxo-6-(1,2-dibromo-2-phenylethyl)pteridine			
	12	4	12
3 2-amino-4-methoxy-6-styryl-7-methoxypteridine			
	25	160	100
4 2-amino-4-methoxy-6-(1,2-dihydroxypropyl)pteridine			
	>200	140	110
5 2,4-diamino-6-phenyl pteridine			
	>200	>200	>200
6 2,4-diamino-6-phenyl-7-methylpteridine			
	>200	>200	>200

7 2,4-diamino-6-(4-tolyl)pteridine



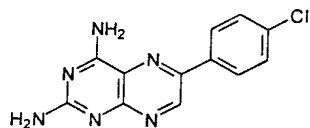
>200 110 25

8 2,4-diamino-6-(4-methoxyphenyl)pteridine



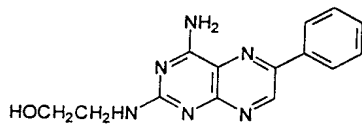
>150 125 22

9 2,4-diamino-6-(4-chlorophenyl)pteridine



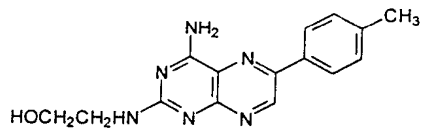
125 110 110

10 2-hydroxyethylamino-4-amino-6-phenylpteridine



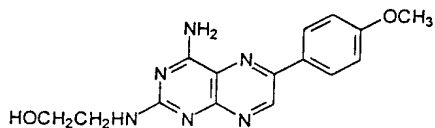
125 125 125

11 2-hydroxyethylamino-4-amino-6-(4-tolyl)pteridine



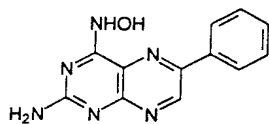
>200 200 160

12 2-hydroxyethylamino-4-amino-6-(4-methoxyphenyl)pteridine



>200 >200 160

13 2-amino-4-hydroxylamino-6-phenylpteridine



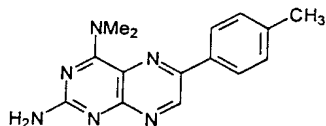
140 >200 >200

14 2-amino-4-dimethylamino-6-phenylpteridine



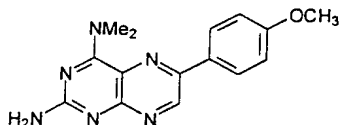
5.0 15 15

15 2-amino-4-dimethylamino-6-(4-tolyl)pteridine



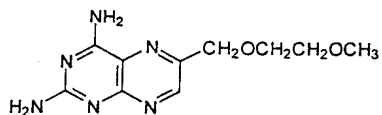
3.6 4.5 3.9

16 2-amino-4-dimethylamino-6-(4-methoxyphenyl)pteridine



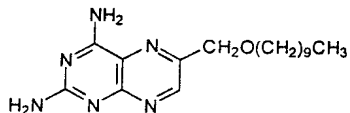
12 7.5 7.5

17 2,4-diamino-6-methoxyethoxymethyl pteridine



>200 180 18

18 2,4-diamino-6-decyloxymethyl pteridine

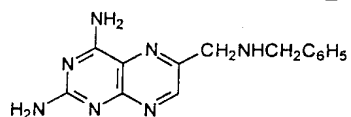


>200 152 100

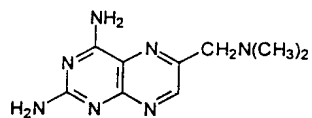
19 2,4-diamino-6-benzylaminomethyl pteridine

50 123 118

24

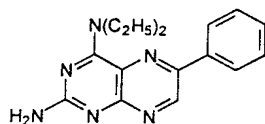


20 2,4-diamino-6-dimethyl aminomethyl pteridine



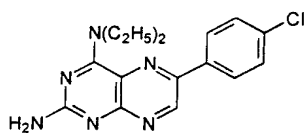
>200 170 100

21 2-amino-4-diethylamino-6-phenylpteridine



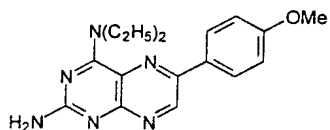
15 80 20

22 2-amino-4-diethylamino-6-(4-chlorophenyl) pteridine



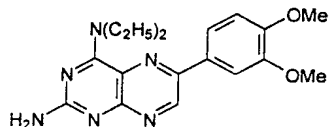
17 50 20

23 2-amino-4-diethylamino-6-(4-methoxyphenyl) pteridine



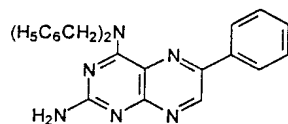
12 20 20

24 2-amino-4-diethylamino-6-(3,4-dimethoxyphenyl) pteridine



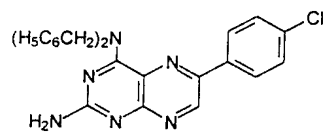
0.08 0.5 0.1

25 2-amino-4-dibenzylamino-6-phenyl pteridine



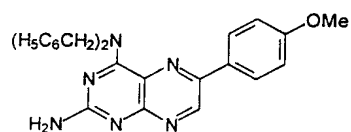
75 25 25

26 2-amino-dibenzylamino-6-(4-chlorophenyl) pteridine



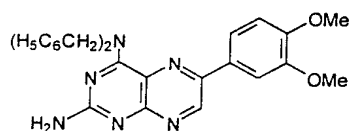
100 75 20

27 2-amino-4-dibenzylamino-6-(4-methoxyphenyl) pteridine



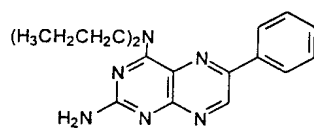
50 20 15

28 2-amino-4-dibenzylamino-6-(3,4-dimethoxyphenyl) pteridine



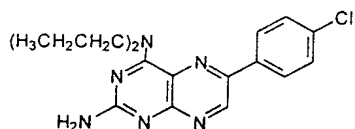
3.5 4.3 3.2

29 2-amino-4-dipropylamino-6-phenylpteridine



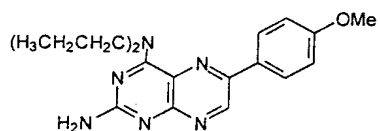
15 25

30 2-amino-4-dipropylamino-6-(4-chlorophenyl)pteridine



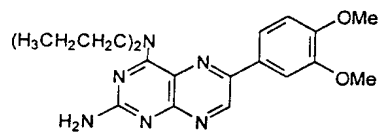
13 9.7 6.7

31 2-amino-4-dipropylamino-6-(4-methoxyphenyl)pteridine



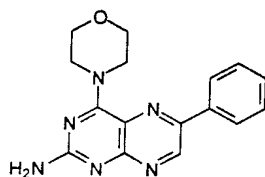
8.5 5.7 2.8

32 2-amino-4-dipropylamino-6-(3,4-dimethoxyphenyl)pteridine



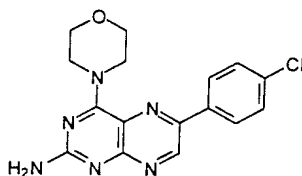
2.2 0.8 0.7

33 2-amino-4-morpholino-6-phenylpteridine



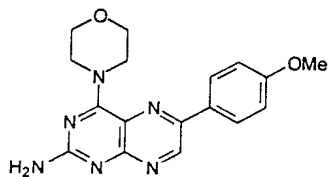
15 100 20

34 2-amino-4-morpholino-6-(4-chlorophenyl)pteridine



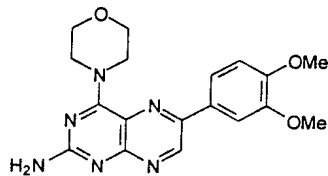
12 20 20

35 2-amino-4-morpholino-6-(4-methoxyphenyl)pteridine



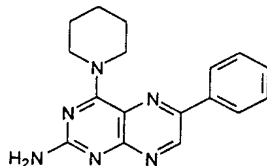
15 125 20

36 2-amino-4-morpholino-6-(3,4-dimethoxyphenyl)pteridine



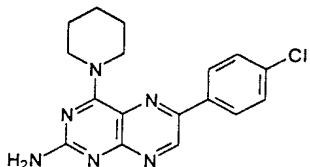
0.1 0.4 0.3

37 2-amino-4-piperidino-6-phenylpteridine



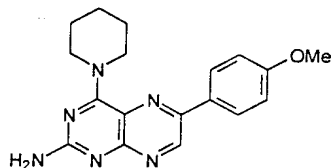
15 80 20

38 2-amino-4-piperidino-6-(4-chlorophenyl)pteridine



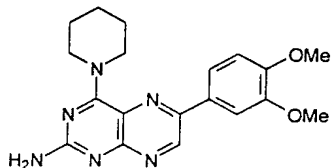
15 70 25

39 2-amino-4-piperidino-6-(4-methoxyphenyl)pteridine



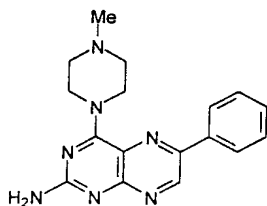
4.6 14 12

40 2-amino-4-piperidino-6-(3,4-dimethoxyphenyl)pteridine



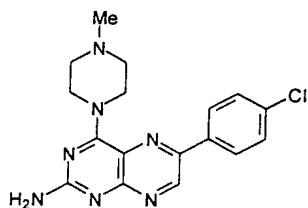
0.4 0.5 0.4

41 2-amino-4-N-methylpiperazino-6-phenylpteridine



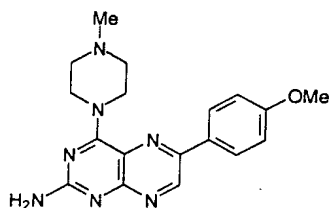
15 40 25

- 42 2-amino-4-N-methylpiperazino-6-(4-chlorophenyl)pteridine



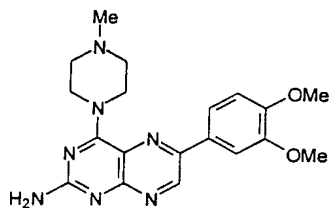
4.8 11 4.2

- 43 2-amino-4-N-methylpiperazino-6-(4-methoxyphenyl)pteridine



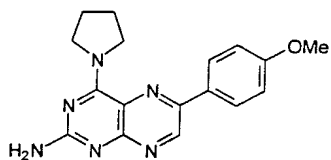
12.3 11.7 4.2

- 44 2-amino-4-methylpiperazino-6-(3,4-dimethoxyphenyl)pteridine



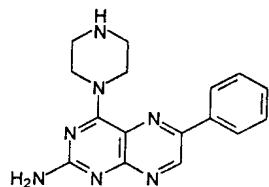
2 7.3 6.5

- 45 2-amino-4-cyclopentylamino-6-(4-methoxyphenyl)pteridine



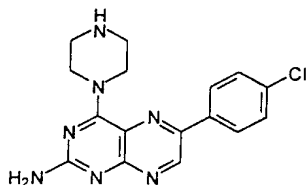
22 17 3.7

- 46 2-amino-4-piperazino-6-phenylpteridine



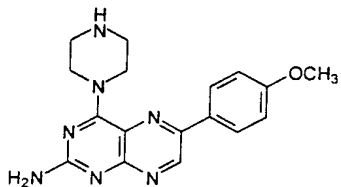
12 11.7 5

- 47 2-amino-4-piperazino-6-(4-chlorophenyl)pteridine



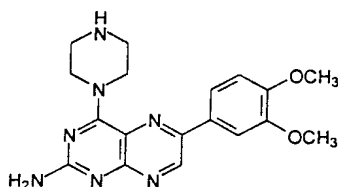
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- 48 2-amino-4-piperazino-6-(4-methoxyphenyl)pteridine



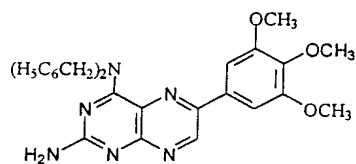
10.5	14	4.5
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- 49 2-amino-4-piperazino-6-(3,4-dimethoxyphenyl)pteridine



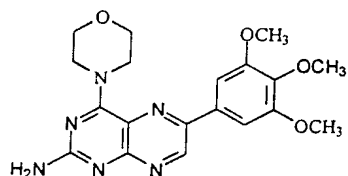
2.8 14.8 4.5

- 50 2-amino-4-dibenzylamino-6-(3,4,5-trimethoxyphenyl)pteridine



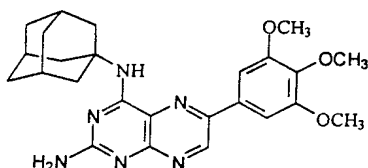
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51 2-amino-4-morpholino-6-(3,4,5-trimethoxyphenyl)pteridine



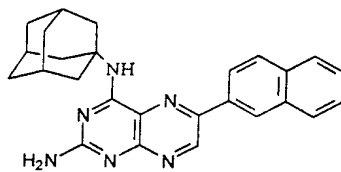
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52 2-amino-4-adamantyl-6-(3,4,5-trimethoxyphenyl)pteridine



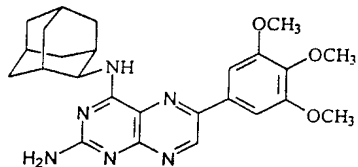
++ ++ ++

53 2-amino-4-adamantyl-6-naftylpteridine



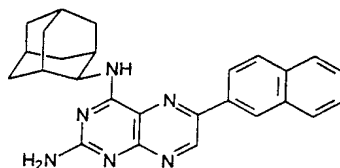
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54 2-amino-4-adamantyl-6-(3,4,5-trimethoxyphenyl)pteridine



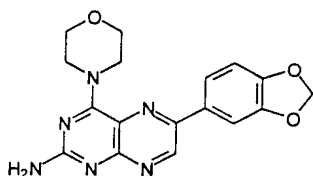
+++ ++ ++

55 2-amino-4-adamantyl-6-naftylpteridine



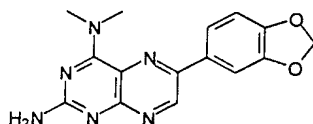
++ ++ ++

- 56 2-amino-4-morpholino-6-(3,4-formylidene-3,4-dihydroxyphenyl)pteridine



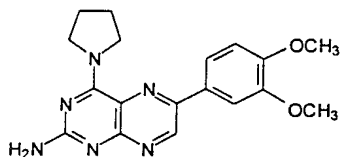
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- 57 2-amino-4-dimethylamino-6-(3,4-formylidene-3,4-dihydroxyphenyl)pteridine



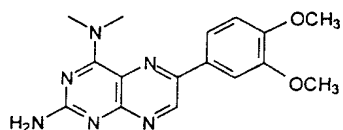
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- 58 2-amino-4-cyclopentylamino-6-(3,4-dimethoxyphenyl)pteridine



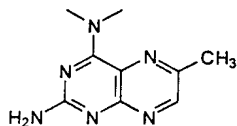
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- 59 2-amino-4-dimethylamino-6-(3,4-dimethoxyphenyl)pteridine



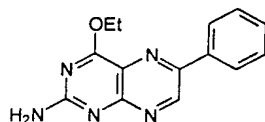
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- 60 2-amino-4-dimethylamino-6-methylpteridine



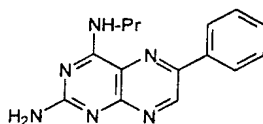
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61 2-amino-4-ethoxy-6-phenylpteridine



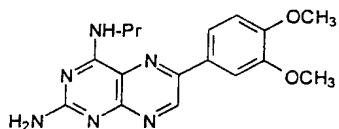
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62 2-amino-4-propylamino-6-phenylpteridine



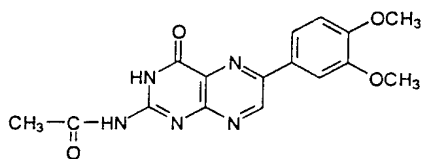
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63 2-amino-4-propylamino-6-(3,4-dimethoxyphenyl)pteridine



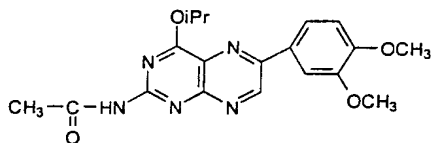
+++ ND ND

64 2-acetamido-4-hydroxy-6-(3,4-dimethoxyphenyl)pteridine



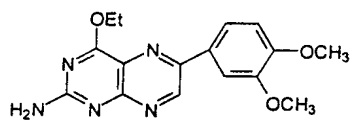
+ 0 +

65 2-acetamido-4-i-propoxy-6-(3,4-dimethoxyphenyl)pteridine



++ + +

66 2-amino-4-ethoxy-6-(3,4-dimethoxyphenyl)pteridine

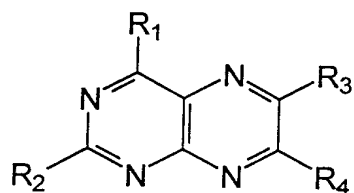


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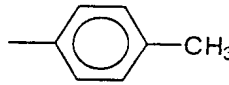
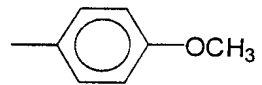
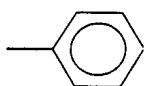
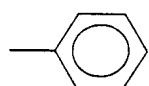
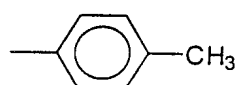
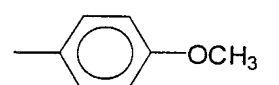
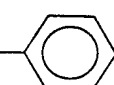
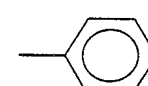
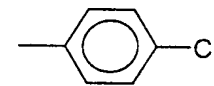
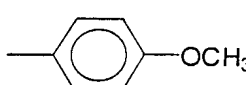
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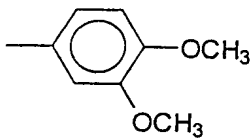
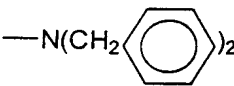
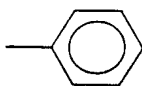
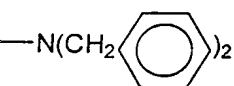
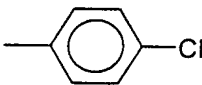
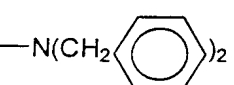
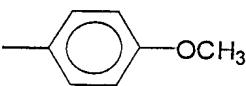
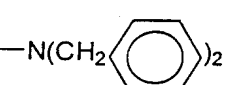
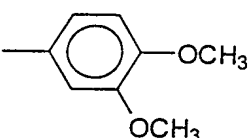
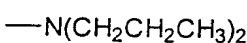
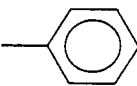
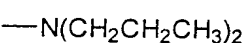
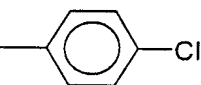
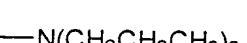
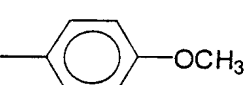
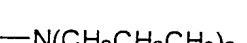
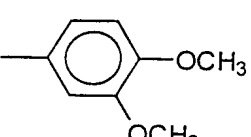
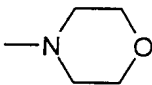
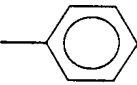
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Table I

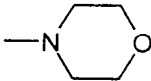
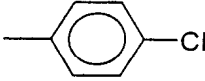
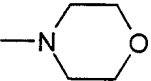
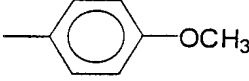
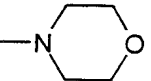
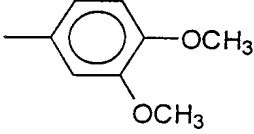
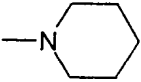
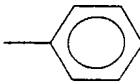
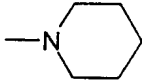
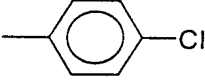
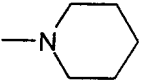
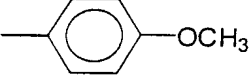
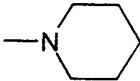
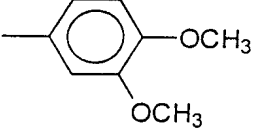
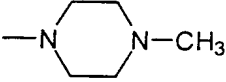
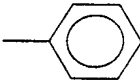
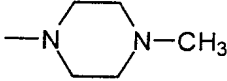

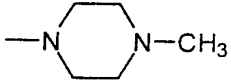
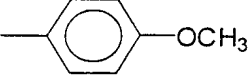


n°	R_1	R_2	R_3	R_4
1	OC_5H_{11}	NH_2	$-CH=CH-$	H
2	OC_5H_{11}	NH_2	$-CHBr-CHBr-$	H
3	OC_5H_{11}	NH_2	$-CH=CH-$	OCH_3
4	OCH_3	NH_2	$-CHOH-CHOH-CH_3$	H
5	NH_2	NH_2		H
6	NMe_2	NH_2		CH_3
7	NH_2	NH_2	$-CH_3$	H
8	NH_2	NH_2	$-OCH_3$	H
9	NH_2	NH_2	$-Cl$	H
10	NH_2	$NHCH_2CH_2OH$		H

11	NH ₂	NHCH ₂ CH ₂ OH		H
12	NH ₂	NH-CH ₂ CH ₂ OH		H
13	NHOH	NH ₂		H
14	NMe ₂	NH ₂		H
15	NMe ₂	NH ₂		H
16	NMe ₂	NH ₂		H
17	NH ₂	NH ₂	CH ₂ OCH ₂ CH ₂ OCH ₃	H
18	NH ₂	NH ₂	CH ₂ O(CH ₂) ₉ CH ₃	H
19	NH ₂	NH ₂	CH ₂ NHCH ₂ - 	H
20	NH ₂	NH ₂	CH ₂ N(CH ₃) ₂	H
21	NEt ₂	NH ₂		H
22	NEt ₂	NH ₂		H
23	NEt ₂	NH ₂		H

24	NEt ₂	NH ₂		H
25		NH ₂		H
26		NH ₂		H
27		NH ₂		H
28		NH ₂		H
29		NH ₂		H
30		NH ₂		H
31		NH ₂		H
32		NH ₂		H
33		NH ₂		H

37

34		NH ₂		H
35		NH ₂		H
36		NH ₂		H
37		NH ₂		H
38		NH ₂		H
39		NH ₂		H
40		NH ₂		H
41		NH ₂		H
42		NH ₂		H
43		NH ₂		H

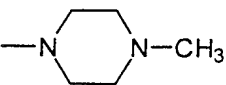
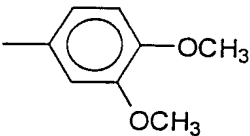
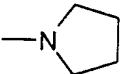
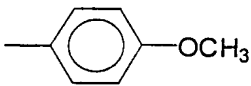
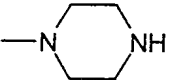
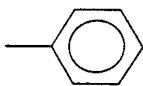
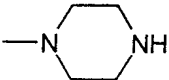
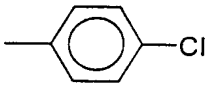
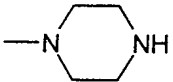
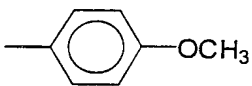
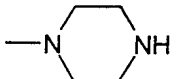
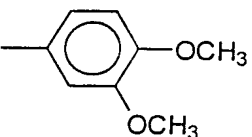
44		NH ₂		H
45		NH ₂		H
46		NH ₂		H
47		NH ₂		H
48		NH ₂		H
49		NH ₂		H

Table II

IC ₅₀ in μ M of pteridine derivative			
Compound n°	MLR	ACD3	aCD28
1	15	110	80
2	12	4	12
3	25	160	100
4	>200	140	110
5	>200	>200	>200
6	>200	>200	>200
7	>200	110	25
8	150	125	22
9	125	110	110
10	125	125	125
11	>200	200	160
12	>200	>200	160
13	140	>200	>200
14	5.0	15	15
15	3.6	4.5	3.9
16	12	7.5	7.5
17	>200	180	18
18	>200	152	100
19	50	123	118
20	>200	170	100
21	15	80	20
22	17	50	20
23	12	20	20
24	0.08	0.5	0.1
25	75	25	25

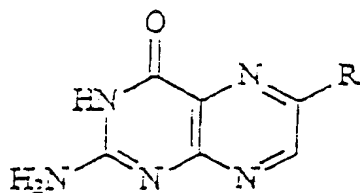
		40	
26	100	75	20
27	50	20	15
28	3.5	4.3	3.2
29	15	25	25
30	13	9.7	6.7
31	8.5	5.7	2.8
32	2.2	0.8	0.7
33	15	100	20
34	12	20	20
35	15	125	20
36	0.1	0.4	0.3
37	15	80	20
38	15	70	25
39	4.6	14	12
40	0.4	0.5	0.4
41	15	40	25
42	4.8	11	4.2
43	12.3	11.7	4.2
44	2	7.3	6.5
45	22	17	3.7
46	12	11.7	5
47	2	20	15
48	10.5	14	4.5
49	2.8	14.8	4.5

Table III

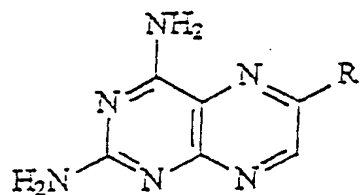
I.S.	IC50		
	Immunosuppressant		
	MLR	aCD3	aCD28
CyA	20 nM	50 nM	N.S.
FK506	1 nM	1 nM	N.S.
Rapamycin	1 nM	1 nM	1 nM
Leflunomide	25 μ M	15 μ M	20 μ M
Mofetil	<0.5 μ M	50 nM	50 nM
MTX	10 μ M	>200 μ M	> 200 μ M
5-FU		50 μ M	17 μ M

N.S. = not suppressive even not in the highest
Concentration

Table IV



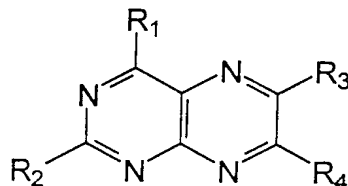
R	MLR	aCD3	aCD23
CH ₂ OOCOCH ₃	0	0	0
CH ₂ OC ₂ H ₅	0	0	0
CH ₂ NHCH ₃	0	0	0
CH ₂ N(CH ₃) ₂	0	0	0
CH ₂ S CH ₃	0	0	0



R	MLR	aCD3	aCD23
CH ₂ S CH ₃	0	0	0
CH ₂ O CH ₂ (CH ₃) ₂	0	0	0
CH ₂ O CH ₃	0	0	0
CH ₂ O CH ₂ CH ₂ CH ₃	0	0	0
CH ₂ NH CO CH (CH ₃) ₂	0	0	0

CLAIMS

1. Pharmaceutical composition for the treatment of autoimmune disorders or for the treatment and/or prevention of transplant-rejections and/or for the treatment of inflammatory diseases comprising as an active principle at least one pteridine derivative having the general formula:



(I)

wherein:

R₁ and R₂ are independently amino, hydroxylamino, (mono- or di)C₁₋₇alkylamino, (mono- or di)C₁₋₇alkyloxyamino, (mono- or di)arylamino, (mono- or di)C₃₋₁₀cycloalkylamino, (mono- or di)hydroxyC₁₋₇alkylamino, (mono- or di)C₁₋₄alkyl-arylamino, mercaptoC₁₋₇alkyl, C₁₋₇alkyloxy or a saturated or unsaturated heterocyclic compound containing at least one nitrogen and optionally substituted by one or more C₁₋₄alkyl, hydroxyC₁₋₄alkyl, C₁₋₄alkyloxy, halo, hydroxy, hydroxycarbonyl, C₁₋₄alkyloxycarbonyl,

R₃ is an unsubstituted, monosubstituted or disubstituted aryl group (wherein the substituent may be, but not limited to, halogen, C₁₋₄alkoxy, C₁₋₄alkyl), aryl group bond to the pteridine ring via a saturated or unsaturated aliphatic spacer which may be halogenated or hydroxylated, aliphatic substituent which may contain ether function, alcohol function, substituted or unsubstituted amino functions or C₁₋₄alkyloxy,

R₄ is hydrogen, alkyl, alkoxy, substituted or unsubstituted aryl

or a pharmaceutical acceptable addition salt or a stereochemical isomeric form thereof.

5 2. Pharmaceutical composition according to claim 1, wherein R₁ is di(C₁₋₄alkyl)amino, morpholinyl, piperidinyl, piperazinyl, C₁₋₄alkylpiperazinyl, pyrrolidinyl or benzylamine.

10 3. Pharmaceutical composition according to claim 1 or 2, wherein R₂ is ammonium, hydroxyammonium, (mono- or di)hydroxylC₁₋₇alkylammonium.

15 4. Pharmaceutical composition according to claim 1, 2 or 3, wherein R₃ is benzyl, phenyl, styryl, phenyl-(C₁₋₄)alkyloxy, phenyl(C₁₋₄alkyl) optionally substituted by one or more C₁₋₄alkyl or halo.

5. Pharmaceutical composition according to claims 1-4, wherein R₄ is hydrogen or C₁₋₄alkyl.

20 6. Pharmaceutical composition according to claims 1-5, wherein the pteridine derivative is a compound chosen from the group comprising:

- 1 2-amino-4-pentoxy-6-styrylpteridine
- 2 2-amino-4-n-pentoxy-6-(1,2-dibromo-2-phenyl ethyl)pteridine
- 3 2-amino-4-methoxy-6-styryl-7-methoxypteridine
- 25 4 2-amino-4-methoxy-6-(1,2-dihydroxypropyl)pteridine
- 5 2,4-diamino-6-phenyl pteridine
- 6 2,4-diamino-6-phenyl-7-methylpteridine
- 7 2,4-diamino-6-(4-tolyl)pteridine
- 8 2,4-diamino-6-(4-methoxyphenyl)pteridine
- 30 9 2,4-diamino-6-(4-chlorophenyl)pteridine
- 10 2-hydroxyethylamino-4-amino-6-phenylpteridine
- 11 2-hydroxyethylamino-4-amino-6-(4-tolyl)pteridine
- 12 2-hydroxyethylamino-4-amino-6-(4-methoxyphenyl)pteridine
- 35 13 2-amino-4-hydroxylamino-6-phenylpteridine
- 14 2-amino-4-dimethylamino-6-phenylpteridine

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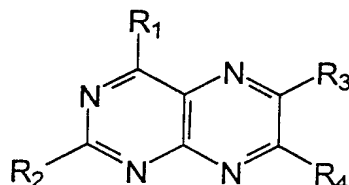
- 15 2-amino-4-dimethylamino-6-(4-tolyl)pteridine
16 2-amino-4-dimethylamino-6-(4-methoxyphenyl)pteridine
17 2,4-diamino-6-methoxyethoxymethyl pteridine
18 2,4-diamino-6-decycloxyethyl pteridine
5 19 2,4-diamino-6-benzylaminomethyl pteridine
20 2,4-diamino-6-dimethyl aminomethyl pteridine
21 2-amino-4-diethylamino-6-phenylpteridine
22 2-amino-4-diethylamino-6-(4-chlorophenyl) pteridine
23 2-amino-4-diethylamino-6-(4-methoxyphenyl) pteridine
10 24 2-amino-4-diethylamino-6-(3,4-dimethoxyphenyl)-
pteridine
25 2-amino-4-dibenzylamino-6-phenyl pteridine
26 2-amino-dibenzylamino-6-(4-chlorophenyl) pteridine
27 2-amino-4-dibenzylamino-6-(4-methoxyphenyl)pteridine
15 28 2-amino-4-dibenzylamino-6-(3,4-dimethoxyphenyl)-
pteridine
29 2-amino-4-dipropylamino-6-phenylpteridine
30 2-amino-4-dipropylamino-6-(4-chlorophenyl)pteridine
31 2-amino-4-dipropylamino-6-(4-methoxyphenyl)pteridine
20 32 2-amino-4-dipropylamino-6-(3,4-dimethoxyphenyl)-
pteridine
33 2-amino-4-morpholino-6-phenylpteridine
34 2-amino-4-morpholino-6-(4-chlorophenyl)pteridine
35 2-amino-4-morpholino-6-(4-methoxyphenyl)pteridine
25 36 2-amino-4-morpholino-6-(3,4-dimethoxyphenyl)pteridine
37 2-amino-4-piperidino-6-phenylpteridine
38 2-amino-4-piperidino-6-(4-chlorophenyl)pteridine
39 2-amino-4-piperidino-6-(4-methoxyphenyl)pteridine
40 2-amino-4-piperidino-6-(3,4-dimethoxyphenyl)pteridine
30 41 2-amino-4-N-methylpiperazino-6-phenylpteridine
42 2-amino-4-N-methylpiperazino-6-(4-chlorophenyl)-
pteridine
43 2-amino-4-N-methylpiperazino-6-(4-methoxyphenyl)-
pteridine
35 44 2-amino-4-methylpiperazino-6-(3,4-dimethoxy-
phenyl)pteridine

- 45 2-amino-4-cyclopentylamino-6-(4-methoxyphenyl)-
pteridine
- 46 2-amino-4-piperazino-6-phenylpteridine
- 5 47 2-amino-4-piperazino-6-(4-chlorophenyl)pteridine
- 48 2-amino-4-piperazino-6-(4-methoxyphenyl)pteridine
- 49 2-amino-4-piperazino-6-(3,4-dimethoxyphenyl)pteridine
- 50 2-amino-4-dibenzylamino-6-(3,4,5-trimethoxyphenyl)-
pteridine
- 10 51 2-amino-4-morpholino-6-(3,4,5-trimethoxyphenyl)-
pteridine
- 52 2-amino-4-adamantyl-6-(3,4,5-trimethoxyphenyl)-
pteridine
- 53 2-amino-4-adamantyl-6-naftylpteridine
- 15 54 2-amino-4-adamantyl-6-(3,4,5-trimethoxyphenyl)-
pteridine
- 55 2-amino-4-adamantyl-6-naftylpteridine
- 56 2-amino-4-morpholino-6-(3,4-formylidene-3,4-
dihydroxyphenyl)pteridine
- 20 57 2-amino-4-dimethylamino-6-(3,4-formylidene-3,4-
dihydroxyphenyl)pteridine
- 58 2-amino-4-cyclopentylamino-6-(3,4-dimethoxyphenyl)-
pteridine
- 59 2-amino-4-dimethylamino-6-(3,4-dimethoxyphenyl)-
pteridine
- 25 60 2-amino-4-dimethylamino-6-methylpteridine
- 61 2-amino-4-ethoxy-6-phenylpteridine
- 62 2-amino-4-propylamino-6-phenylpteridine
- 63 2-amino-4-propylamino-6-(3,4-dimethoxyphenyl)-
pteridine
- 30 64 2-acetamido-4-hydroxy-6-(3,4-dimethoxyphenyl)-
pteridine
- 65 2-acetamido-4-i-propoxy-6-(3,4-dimethoxyphenyl)-
pteridine
- 35 66 2-amino-4-ethoxy-6-(3,4-dimethoxyphenyl)pteridine

7. Pharmaceutical composition according to

claims 1-6, further comprising one or more other immunosuppressants chosen from the group comprising CyA, FK506, rapamycin (RPM), leflunomide, mofetil, MTX or 5-FU.

5 8. Compound having the general formula:



(I)

10 wherein:

R₁ and R₂ are independently amino, hydroxylamino, (mono- or di)C₁₋₇alkylamino, (mono- or di)C₁₋₇alkyloxyamino, (mono- or di)arylamino, (mono- or di)C₃₋₁₀cycloalkylamino, (mono- or di)hydroxyC₁₋₇alkylamino, (mono- or di)C₁₋₄alkyl-
15 arylamino, mercaptoC₁₋₇alkyl, C₁₋₇alkyloxy or a saturated or unsaturated heterocyclic compound containing at least one nitrogen and optionally substituted by one or more C₁₋₄alkyl, hydroxyC₁₋₄alkyl, C₁₋₄alkyloxy, halo, hydroxy, hydroxycarbonyl, C₁₋₄alkyloxycarbonyl,

20 R₃ is an unsubstituted, monosubstituted or disubstituted aryl group (wherein the substituent may be, but not limited to, halogen, C₁₋₄alkoxy, C₁₋₄alkyl), aryl group bond to the pteridine ring via a saturated or unsaturated aliphatic spacer which may be halogenated or
25 hydroxylated, aliphatic substituent which may contain ether function, alcohol function, substituted or unsubstituted amino functions or C₁₋₄alkyloxy,

R₄ is hydrogen, alkyl, alkoxy, substituted or unsubstituted aryl

30 or a pharmaceutical acceptable addition salt or a stereochemical isomeric form thereof.

9. Compound according to claim 8, wherein R₁ is

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di(C₁₋₄alkyl)amino, morpholinyl, piperidinyl, piperazinyl, C₁₋₄alkylpiperazinyl, pyrrolidinyl or benzylamine.

10. Compound according to claim 8 or 9, wherein R₂ is ammonium, hydroxyammonium, (mono- or di)hydroxyl-
5 C₁₋₇alkylammonium.

11. Compound according to claims 8-10, wherein R₃ is benzyl, phenyl, styryl, phenyl(C₁₋₄)alkyloxy, phenyl-(C₁₋₄alkyl) optionally substituted by one or more C₁₋₄alkyl or halo.

10 12. Compound according to claims 8-11, wherein R₄ is hydrogen or C₁₋₄alkyl.

13. Compound having the formula:

- | | |
|----|--|
| 1 | 2-amino-4-pentoxo-6-styrylpteridine |
| 2 | 2-amino-4-n-pentoxo-6-(1,2-dibromo-2-phenylethyl)- |
| 15 | pteridine |
| 3 | 2-amino-4-methoxy-6-styryl-7-methoxypteridine |
| 6 | 2,4-diamino-6-phenyl-7-methylpteridine |
| 14 | 2-amino-4-dimethylamino-6-phenylpteridine |
| 15 | 2-amino-4-dimethylamino-6-(4-tolyl)pteridine |
| 20 | 16 2-amino-4-dimethylamino-6-(4-methoxyphenyl)pteridine. |
| 21 | 2-amino-4-diethylamino-6-phenylpteridine |
| 22 | 2-amino-4-diethylamino-6-(4-chlorophenyl) pteridine |
| 23 | 2-amino-4-diethylamino-6-(4-methoxyphenyl) pteridine |
| 24 | 2-amino-4-diethylamino-6-(3,4-dimethoxyphenyl)- |
| 25 | pteridine |
| 25 | 2-amino-4-dibenzylamino-6-phenyl pteridine |
| 26 | 2-amino-dibenzylamino-6-(4-chlorophenyl) pteridine |
| 27 | 2-amino-4-dibenzylamino-6-(4-methoxyphenyl)pteridine |
| 28 | 2-amino-4-dibenzylamino-6-(3,4-dimethoxyphenyl)- |
| 30 | pteridine |
| 29 | 2-amino-4-dipropylamino-6-phenylpteridine |
| 30 | 2-amino-4-dipropylamino-6-(4-chlorophenyl)pteridine |
| 31 | 2-amino-4-dipropylamino-6-(4-methoxyphenyl)pteridine |
| 32 | 2-amino-4-dipropylamino-6-(3,4-dimethoxyphenyl)- |
| 35 | pteridine |
| 33 | 2-amino-4-morpholino-6-phenylpteridine |

- 34 2-amino-4-morpholino-6-(4-chlorophenyl)pteridine
35 2-amino-4-morpholino-6-(4-methoxyphenyl)pteridine
36 2-amino-4-morpholino-6-(3,4-dimethoxyphenyl)-
pteridine
5 37 2-amino-4-piperidino-6-phenylpteridine
38 2-amino-4-piperidino-6-(4-chlorophenyl)pteridine
39 2-amino-4-piperidino-6-(4-methoxyphenyl)pteridine
40 2-amino-4-piperidino-6-(3,4-dimethoxyphenyl)-
pteridine
10 41 2-amino-4-N-methylpiperazino-6-phenylpteridine
42 2-amino-4-N-methylpiperazino-6-(4-chlorophenyl)-
pteridine
43 2-amino-4-N-methylpiperazino-6-(4-methoxyphenyl)-
pteridine
15 44 2-amino-4-methylpiperazino-6-(3,4-dimethoxyphenyl)-
pteridine
45 2-amino-4-cyclopentylamino-6-(4-methoxyphenyl)-
pteridine
46 2-amino-4-piperazino-6-phenylpteridine
20 47 2-amino-4-piperazino-6-(4-chlorophenyl)pteridine
48 2-amino-4-piperazino-6-(4-methoxyphenyl)pteridine
49 2-amino-4-piperazino-6-(3,4-dimethoxyphenyl)-
pteridine
50 2-amino-4-dibenzylamino-6-(3,4,5-trimethoxyphenyl)-
25 pteridine
51 2-amino-4-morpholino-6-(3,4,5-trimethoxyphenyl)-
pteridine
51 2-amino-4-adamantyl-6-(3,4,5-trimethoxyphenyl)-
pteridine
30 53 2-amino-4-adamantyl-6-naftylpteridine
54 2-amino-4-adamantyl-6-(3,4,5-trimethoxyphenyl)-
pteridine
55 2-amino-4-adamantyl-6-naftylpteridine
56 2-amino-4-morpholino-6-(3,4-formylidene-3,4-
35 dihydroxyphenyl)pteridine
57 2-amino-4-dimethylamino-6-(3,4-formylidene-3,4-

dihydroxyphenyl)pteridine

- 58 2-amino-4-cyclopentylamino-6-(3,4-dimethoxyphenyl)-
pteridine
- 5 58 2-amino-4-dimethylamino-6-(3,4-dimethoxyphenyl)-
pteridine
- 60 2-amino-4-dimethylamino-6-methylpteridine
- 61 2-amino-4-ethoxy-6-phenylpteridine
- 62 2-amino-4-propylamino-6-phenylpteridine
- 63 2-amino-4-propylamino-6-(3,4-dimethoxyphenyl)-
10 pteridine
- 64 2-acetamido-4-hydroxy-6-(3,4-dimethoxyphenyl)-
pteridine
- 65 2-acetamido-4-i-propoxy-6-(3,4-dimethoxyphenyl)-
pteridine
- 15 66 2-amino-4-ethoxy-6-(3,4-dimethoxyphenyl)pteridine.

14. Use of a compound according to claims 8-13
as a medicament.

15. Use of a compound according to claims 8-13,
for the manufacture of a medicament for the treatment of
20 autoimmuno disorders.

16. Use of a compound according to claims 8-13,
for the manufacture of a medicament for the prevention of
transplant-rejections.

17. Use of a compound according to claims 8-13,
25 for the manufacture of a medicament for the treatment of
inflammatory diseases.

18. Method for selecting immunosuppressive
agents by a combination of at least three test systems
based on MLR, aCD3 and aCD28.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 99/10320

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D475/04 C07D475/08 A61K31/519 A61P35/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D A61K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	EP 0 362 645 A (THOMAE GMBH DR K) 11 April 1990 (1990-04-11) claims	1-18
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	— — — — — -/-	



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

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- "O" document referring to an oral disclosure, use, exhibition or other means
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"&" document member of the same patent family

Date of the actual completion of the international search

23 March 2000

Date of mailing of the international search report

03/04/2000

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Chouly, J

INTERNATIONAL SEARCH REPORT

International Application No

PCT/EP 99/10320

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
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information on patent family members

Inter: nal Application No

PCT/EP 99/10320

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INTERNATIONAL SEARCH REPORT

information on patent family members

International Application No

PCT/EP 99/10320

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
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